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FILE 'USPAT' ENTERED AT 13:38:54 ON 25 JUN 1999
           U.S. PATENT TEXT FILE
    THE WEEKLY PATENT TEXT AND IMAGE DATA IS CURRENT
    THROUGH June 22,1999.
=> file uspat usocr
FILE 'USPAT' ENTERED AT 13:39:18 ON 25 JUN 1999
                  PATENT TEXT
   THE WEEKLY PATENT TEXT AND IMAGE DATA IS CURRENT
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FILE 'USOCR' ENTERED AT 13:39:18 ON 25 JUN 1999
=> s (514/314)/ccls
FILE 'USPAT'
L1 709 (514/314)/CCLS
FILE 'USOCR'
         709 (514/314)/CCLS
TOTAL FOR ALL FILES
L3 1418 (514/314)/CCLS
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Page 2

=> s L3 and indol?

FILE 'USPAT'

24511 INDOL?

L4 328 L1 AND INDOL?

FILE 'USOCR'

1253 INDOL?

L5 0 L2 AND INDOL?

TOTAL FOR ALL FILES

L6 328 L3 AND INDOL?

=> s L6 and glyoxylamid?

FILE 'USPAT'

80 GLYOXYLAMID?

L7 0 L4 AND GLYOXYLAMID?

FILE 'USOCR'

5 GLYOXYLAMID?

L8 0 L5 AND GLYOXYLAMID?

TOTAL FOR ALL FILES

L9 0 L6 AND GLYOXYLAMID?

=> s (514/339)/ccls

FILE 'USPAT'

L10 394 (514/339)/CCLS

FILE 'USOCR'

L11 394 (514/339)/CCLS

TOTAL FOR ALL FILES

L12 788 (514/339)/CCLS

=> s L12 and indol?

FILE 'USPAT'

24511 INDOL?

L13 268 L10 AND INDOL?

FILE 'USOCR'

1253 INDOL?

L14 4 L11 AND INDOL?

TOTAL FOR ALL FILES

L15 272 L12 AND INDOL?

=> s L15 and glyoxylamid?

FILE 'USPAT'

80 GLYOXYLAMID?

L16 1 L13 AND GLYOXYLAMID?

FILE 'USOCR'

5 GLYOXYLAMID?

L17 0 L14 AND GLYOXYLAMID?

TOTAL FOR ALL FILES

L18 1 L15 AND GLYOXYLAMID?

=> d L18 ti in pd ab ccls

US PAT NO: 5,380,723 [IMAGE AVAILABLE] L18: 1 of 1

TITLE: Indole derivatives

INVENTOR: Toshihiro Takahashi, Saitama, Japan

Hitoshi Inoue, Saitama, Japan Masato Horigome, Tokyo, Japan Kenichi Momose, Saitama, Japan Masanori Sugita, Saitama, Japan Kouichi Katsuyama, Saitama, Japan Chikako Suzuki, Saitama, Japan Shinji Nagai, Saitama, Japan Masao Nagase, Saitama, Japan Koichi Nakamaru, Saitama, Japan

DATE ISSUED: Jan. 10, 1995

ABSTRACT:

Disclosed are **indole** derivatives of formula (I) ##STR1## wherein X and Y each independently represent H or --CH.sub.2 CH.sub.2 R; R represents pyridyl or substituted amino of NR.sub.1 R.sub.2;

R.sub.1 represents H or C.sub.1 -C.sub.6 alkyl;

R.sub.2 represents 2-(3-indoly1)ethyl or aralkyl; or

R.sub.1 and R.sub.2 together with the nitrogen atom to which they are attached may form an N-containing 5 to 6 membered hetero ring, an N, O-containing hetero ring, which hetero ring may be fused with benzene;

n is an integer of 4 to 8;

with the proviso that X and Y both do not represent H or piperizinoethyl when n is 4, or pharmaceutically acceptable acid addition salts thereof. They are useful as an antiulcer agent.

Page 4

US-CL-CURRENT: 514/235.8, 310, 323, 339, 414; 544/143; 546/148, 165, 201, 277.4; 548/455

=> s (514/419)/ccls

FILE 'USPAT'

455 (514/419)/CCLS L19

FILE 'USOCR'

455 (514/419)/CCLS L20

TOTAL FOR ALL FILES

910 (514/419)/CCLS L21

=> s L21 and indol?

FILE 'USPAT'

24511 INDOL?

312 L19 AND INDOL? L22

FILE 'USOCR'

1253 INDOL?

3 L20 AND INDOL? L23

TOTAL FOR ALL FILES

315 L21 AND INDOL? L24

=> s L24 and glyoxylamid?

FILE 'USPAT'

80 GLYOXYLAMID?

L25 4 L22 AND GLYOXYLAMID?

FILE 'USOCR'

5 GLYOXYLAMID?

L26 0 L23 AND GLYOXYLAMID?

TOTAL FOR ALL FILES

L27 4 L24 AND GLYOXYLAMID?

 \Rightarrow d L27 1-4 ti in pd ab ccls

US PAT NO: 5,733,923 [IMAGE AVAILABLE] L27: 1 of 4

1H-indole-3-glyoxylamide sPLA.sub.2 inhibitors TITLE:

INVENTOR: Nicholas J. Bach, Indianapolis, IN

Robert D. Dillard, Zionsville, IN

Susan E. Draheim, Indianapolis, IN

DATE ISSUED: Mar. 31, 1998

ABSTRACT:

A class of novel 1H-indole-3-glyoxylamides is disclosed together with the use of such indole compounds for inhibiting sPLA.sub.2 mediated release of fatty acids for treatment of conditions such as septic shock.

US-CL-CURRENT: 514/419; 548/447, 494

L27: 2 of 4 5,684,034 [IMAGE AVAILABLE] US PAT NO:

1H-indole-3-acetamide sPLA.sub.2 inhibitors TITLE:

Nicholas J. Bach, Indianapolis, IN INVENTOR:

Robert D. Dillard, Zionsville, IN Susan E. Draheim, Indianapolis, IN Robert B. Hermann, Indianapolis, IN Richard W. Schevitz, Indianapolis, IN

DATE ISSUED: Nov. 4, 1997

ABSTRACT:

A class of novel 1-indole-3-acetamides represented by the formula; ##STR1## is disclosed together with the use of such indole compounds for inhibiting sPLA.sub.2 mediated release of fatty acids. US-CL-CURRENT: **514/419**, 418; 548/113, 483, 486, 493, 496

US PAT NO: 5,654,326 [IMAGE AVAILABLE] L27: 3 of 4

1H-indole-3-glyoxylamide SPLA.sub.2 inhibitors TITLE:

INVENTOR:

Nicholas J. Bach, Indianapolis, IN Robert D. Dillard, Zionsville, IN

Susan E. Draheim, Indianapolis, IN

DATE ISSUED: Aug. 5, 1997

ABSTRACT:

A class of novel 1H-indole-3-glyoxylamides is disclosed together with the use of such indole compounds for inhibiting sPLA.sub.2 mediated release of fatty acids for treatment of conditions such as septic shock.

US-CL-CURRENT: 514/419, 381; 544/373; 548/492, 494, 495

L27: 4 of 4 5,578,634 [IMAGE AVAILABLE] US PAT NO:

1H-indole-3-acetic acid hydrazide sPLA.sub.2 TITLE:

inhibitors

Nicholas J. Bach, Indianapolis, IN INVENTOR:

Robert D. Dillard, Zionsville, IN Susan E. Draheim, Indianapolis, IN Robert B. Hermann, Indianapolis, IN Richard W. Schevitz, Indianapolis, IN

DATE ISSUED: Nov. 26, 1996

ABSTRACT:

A class of novel 1H-indole-3-acetic acid hydrazides is disclosed together with the use of such indole compounds for inhibiting sPLA.sub.2 mediated release of fatty acids (e.g., arachidonic acid) for treatment of conditions such as septic shock.

US-CL-CURRENT: 514/419, 92, 362, 381; 548/135, 253, 414, 494

=> s (546/168)/ccls

FILE 'USPAT'

L28 306 (546/168)/CCLS

FILE 'USOCR'

L29 306 (546/168)/CCLS

TOTAL FOR ALL FILES

L30 612 (546/168)/CCLS

=> s L30 and indol?

FILE 'USPAT'

24511 INDOL?

L31 96 L28 AND INDOL?

FILE 'USOCR'

1253 INDOL?

L32 4 L29 AND INDOL?

TOTAL FOR ALL FILES

L33 100 L30 AND INDOL?

=> s L33 and quinolin?

FILE 'USPAT'

24713 QUINOLIN?

L34 59 L31 AND QUINOLIN?

FILE 'USOCR'

2148 QUINOLIN?

L35 4 L32 AND QUINOLIN?

TOTAL FOR ALL FILES

L36 63 L33 AND QUINOLIN?

=> s L36 and glyoxylamid? .

FILE 'USPAT' 80 GLYOXYLAMID? 0 L34 AND GLYOXYLAMID? L37 FILE 'USOCR' 5 GLYOXYLAMID? 0 L35 AND GLYOXYLAMID? TOTAL FOR ALL FILES 0 L36 AND GLYOXYLAMID? L39 => s (546/278.1)FILE 'USPAT' *WARNING* - FIELD CODE NOT VALID '278.1' L40 0 (546/278.1) FILE 'USOCR' *WARNING* - FIELD CODE NOT VALID '278.1' 0 (546/278.1) L41 TOTAL FOR ALL FILES 0 (546/278.1) L42 => s (546/278.1)/cclsFILE 'USPAT' 157 (546/278.1)/CCLS L43 FILE 'USOCR' 157 (546/278.1)/CCLS TOTAL FOR ALL FILES 314 (546/278.1)/CCLS => s L45 and indol? FILE 'USPAT' 24511 INDOL? L46 97 L43 AND INDOL? FILE 'USOCR'

1253 INDOL?

TOTAL FOR ALL FILES

22 L44 AND INDOL?

L47

L48 119 L45 AND INDOL?

=> s L48 and glyoxylamid?

FILE 'USPAT'

80 GLYOXYLAMID?

L49 0 L46 AND GLYOXYLAMID?

FILE 'USOCR'

5 GLYOXYLAMID?

L50 1 L47 AND GLYOXYLAMID?

TOTAL FOR ALL FILES

L51 1 L48 AND GLYOXYLAMID?

=> d 151 ti in pd ab ccls

US PAT NO: OCR DATA 3,801,594 [IMAGE AVAILABLE] L51: 1 of 1

TITLE: SUBSTI*

INVENTOR: NAME MAY BE IN MISC FIELD, CITY MAY BE IN MISC FIELD

DATE ISSUED: Apr. 2, 1974

ABSTRACT:

ABSTRACT OF THE IIISCLOSURE This invention describes new substituted indole compounds. Intermediate 3-halodicarbonyl indoles are also described. The preparation of *e final products from the . 3-halodicarbonyl indoles wherein the 3-position contains a disubstituted g-aminoethyl group is described. The compounds have utility as central nervous system depressants, analgesics, anti-inflammatory agents, and illuretic agents.

US-CL-CURRENT: 540/582; 514/869; 544/144; 546/112, 277.4, **278.1**; 548/468, 492, 493, 504

=> s (548?491)/ccls

FILE 'USPAT'

'?' TRUNCATION SYMBOL NOT VALID WITHIN '548?491'

FILE 'USOCR'

'?' TRUNCATION SYMBOL NOT VALID WITHIN '548?491'
YOU HAVE RECEIVED THIS ERROR MESSAGE 2 CONSECUTIVE TIMES
The truncation symbol ? may be used only at the end of a search
term. To specify a variable character within a word use '!', e.g.,
'wom!n' to search for both 'woman' and 'women'. Enter "HELP
TRUNCATION" at an arrow prompt (=>) for more information.

IF YOU REQUIRE FURTHER HELP, PLEASE CONTACT YOUR LOCAL HELP DESK
=> s (548/491)/ccls

FILE 'USPAT'

L52 341 (548/491)/CCLS

FILE 'USOCR'

L53 341 (548/491)/CCLS

TOTAL FOR ALL FILES

L54 682 (548/491)/CCLS

=> s L54 and indol?

FILE 'USPAT'

24511 INDOL?

L55 210 L52 AND INDOL?

FILE 'USOCR'

1253 INDOL?

L56 27 L53 AND INDOL?

TOTAL FOR ALL FILES

L57 237 L54 AND INDOL?

=> s L57 and glyoxylamid?

FILE 'USPAT'

80 GLYOXYLAMID?

L58 0 L55 AND GLYOXYLAMID?

FILE 'USOCR'

5 GLYOXYLAMID?

L59 0 L56 AND GLYOXYLAMID?

TOTAL FOR ALL FILES

L60 0 L57 AND GLYOXYLAMID?

=> s (548/493)/ccls

FILE 'USPAT'

L61 184 (548/493)/CCLS

FILE 'USOCR'

L62 184 (548/493)/CCLS

TOTAL FOR ALL FILES

L63 368 (548/493)/CCLS

=> s L63 and glyoxylamid?

FILE 'USPAT'

80 GLYOXYLAMID?

L64 2 L61 AND GLYOXYLAMID?

FILE 'USOCR'

5 GLYOXYLAMID?

L65 2 L62 AND GLYOXYLAMID?

TOTAL FOR ALL FILES

L66 4 L63 AND GLYOXYLAMID?

=> d L66 1-4 ti in pd ab ccls

US PAT NO: 5,684,034 [IMAGE AVAILABLE] L66: 1 of 4

TITLE: 1H-indole-3-acetamide sPLA.sub.2 inhibitors

INVENTOR: Nicholas J. Bach, Indianapolis, IN

Robert D. Dillard, Zionsville, IN Susan E. Draheim, Indianapolis, IN Robert B. Hermann, Indianapolis, IN Richard W. Schevitz, Indianapolis, IN

DATE ISSUED: Nov. 4, 1997

ABSTRACT:

A class of novel 1-indole-3-acetamides represented by the formula; ##STR1## is disclosed together with the use of such indole compounds for inhibiting sPLA.sub.2 mediated release of fatty acids.
US-CL-CURRENT: 514/419, 418; 548/113, 483, 486, 493, 496

US PAT NO: 3,915,990 [IMAGE AVAILABLE] L66: 2 of 4

TITLE: Tryptamines

INVENTOR: John R. Smythies, Birmingham, AL

DATE ISSUED: Oct. 28, 1975

ABSTRACT:

Substituted indoles and benzimidazoles having serotonin blocking activity and having the structural formulae #EQU1# and #EQU2# wherein one and only one of Y or Z is --OR.sub.3 and WHEREIN R.sub.1, R.sub.2 and R.sub.3 is H or lower alkyl, and

Wherein R.sub.4 is a phenyl group or phenyl group substituted with lower alkyl, lower alkoxy, halogen, CF.sub.3, NH.sub.2, NO.sub.2, CN, --NH-lower alkyl or ##EQU3## GROUP.

US-CL-CURRENT: 548/506; 514/903, 923; 546/86; 548/309.7, 493, 504,

505, 507

US PAT NO: OCR DATA 3,801,594 [IMAGE AVAILABLE] L66: 3 of 4

TITLE: SUBSTI*

INVENTOR: NAME MAY BE IN MISC FIELD, CITY MAY BE IN MISC FIELD

DATE ISSUED: Apr. 2, 1974

ABSTRACT:

ABSTRACT OF THE IIISCLOSURE This invention describes new substituted indole compounds. Intermediate 3-halodicarbonyl indoles are also described. The preparation of *e final products from the .

3-halodicarbonyl indoles whereiu the 3-position contains a disubstituted g-aminoethyl group is described. The compounds have utility as central nervous system depressants, analgesics, anti-inflammatory agents, and iliuretic agents.

US-CL-CURRENT: 540/582; 514/869; 544/144; 546/112, 277.4, 278.1; 548/468, 492, 493, 504

US PAT NO: OCR DATA 3,686,213 [IMAGE AVAILABLE] L66: 4 of 4

TITLE: TITLE MAY BE IN MISC FIELD

INVENTOR: NAME MAY BE IN MISC FIELD, CITY MAY BE IN MISC FIELD

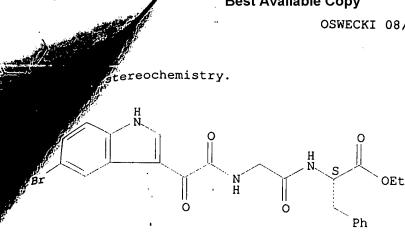
DATE ISSUED: Aug. 22, 1972

ABSTRACT:

15 ta 7 'ken together is morpholino, pyrrolinyl, 1,2,5,6-tetrahyThis invention describes new substituted indole com- dropyridyl, 2-lower alkyl US-CL-CURRENT: 548/504; 514/869; 540/582, 602; 544/144; 546/16, 112, 201, 277.4; 548/468, 493, 507; 560/155

=> log y

U.S. Patent & Trademark Office LOGOFF AT 13:50:18 ON 25 JUN 1999



RN 153694-29-2 HCAPLUS

CN L-Phenylalanine, N-[N-[(5-nitro-1H-indol-3-yl)oxoacetyl]glycyl]-, ethyl ester (9CI) (CA INDEX NAME)

Absolute stereochemistry.

Ι

ANSWER 12 OF 30 HCAPLUS COPYRIGHT 1998 ACS 1994:125105 Document No. 120:125105 Probing the 5-HT3 receptor site using novel indole-3-glyoxylic acid derivatives. Evans, S. M.; Huang, B. S.; Feng, D.; Gall, M.; Tsai, C.; Bariso, C.; Taylor, C. A. (ABOC Health Care Co., Anaquest Inc., Murray Hill, NJ, 07974, USA). Med. Chem. Res., 3(5-6), 386-406 (English) 1993. CODEN: MCREEB. ISSN: 1054-2523. GI

AΒ Novel ester and amide derivs. of indole-3-glyoxylic acid were synthesized and used to probe the 5-HT3 receptor binding site. The structural design of these ligands was based on 1) the rigidity and preferred conformation of the glyoxylic acid fragment, as shown by ab initio geometry optimization using the 3-21G basis set, and 2) the chem. template comprising the 3-dimensional pharmacophore for the 5-HT3 recognition site. The geometrical changes provide ligands which are selective for the 5-HT3 receptor and demonstrate good antiemetic potency. The most potent compd. (I) had a binding affinity of 33 nM and an ED50 of 0.07 mg/kg i.v. in the KATHLEEN FULLER BT/LIBRARY 308-4290

fisplatin-induced emesis assay in ferrets.

2-2 (Mammalian Hormones)

serotoninergic S3 receptor ligand; indoleglyoxylate deriv serotonin receptor

Pharmacophores ΙT

(of serotoninergic S3 receptors)

IT Receptors

IT

IT

GI

RL: BIOL (Biological study)

(serotoninergic S3, indoleglyoxylate derivs. as ligands for)

132797-95-6P . 143137-38-6P 152721-50-1P 152721-51-2P 152721-55-6P 152721-54-5P 152721-52-3P . 152721-53-4P 152721-58-9P 152721-59-0P 152721-56-7P **152721-57-8P**

152721-60-3P

RL: SPN (Synthetic preparation); PREP

(Preparation)

(prepn. and serotoninergic S3 receptor binding of) 51605-33-5P, 4-Chloromethyl-5-methylimidazole hydrochloride 152721-61-4P 152721-62-5P 152721-63-6P 72631-77**-**7P

RL: SPN (Synthetic preparation); PREP (Preparation) (prepn. of)

152721-57-8P

RL: SPN (Synthetic preparation); PREP

(Preparation)

(prepn. and serotoninergic S3 receptor binding of)

152721-57-8 HCAPLUS RN

1H-Indole-3-acetamide, .alpha.-oxo-N-3-pyridinyl- (9CI) (CA INDEX CN NAME)

L24 ANSWER 13 OF 30 HCAPLUS COPYRIGHT 1998 ACS 1991:449510 Document No. 115:49510 Synthesis and antihypertensive activity of some 2-aminobenzimidazole and indole derivatives. Da Settimo, Antonio; Marini, Anna Maria; Primofiore, Giampaolo; Subissi, Alessandro (Ist. Chim. Farm., Univ. Pisa, Pisa, 56100, Italy). Farmaco, 46(2), 357-67 (English) 1991. CODEN: FRMCE8.

Aminobenzimidazole derivs. I [R = H, CH2Ph, Me, CH2C6H4Cl-4, Rl' AB NHCOCOR4, R2R3 = bond, R4 = 2,6-dichloroanilino (throughout); R = H, CH2Ph, Me, CH2C6H4Cl-4, R1R2 = NH, R3 = CH2COR4] and indole derivs. II (R5 = COCOR4, R6, R7 = H, Me, R8 = H, Br, C1, NO2, OMe; R5 = CH2COR4, R6 = R7 = R8 = H) were prepd. and some were tested for antihypertensive activity. Thus, indol-3-ylacetyl chloride condensed with 2,6-dichloroaniline to give II (R5 = CH2COR4, R6 = R7 = R8 = H). None of the compds. tested showed appreciable antihypertensive activity.

KATHLEEN FULLER BT/LIBRARY 308-4290